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25. The method of claim 23, wherein the selected gene encodes a therapeutic protein useful for treating a blood disorder.

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26. The method of claim 25, wherein the therapeutic protein is erythropoietin.

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27. The method of claim 25, wherein the blood disorder is hemophilia.

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28. The method of claim 23 wherein said protein is secreted.

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29. A method of expressing a therapeutically effective amount of a protein in a mammalian subject, said method comprising:

administering into the bloodstream of said subject a pharmaceutical composition which comprises (a) a pharmaceutically acceptable excipient; and (b) recombinant AAV virions comprising a selected gene operably linked to expression control elements that provide for transcription and translation of the selected gene in a desired host cell *in vivo*, whereby said virions transduce cells in said subject, and said selected gene is expressed by the transduced cells at a level which provides for a therapeutic effect in said subject.

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30. The method of claim 29, wherein the pharmaceutical composition is delivered intraarterially.

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31. The method of claim 29, wherein the selected gene encodes a therapeutic protein useful for treating a blood disorder.

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32. The method of claim 31, wherein the therapeutic protein is erythropoietin.